

MAR05-2004-003115

Abstract for an Invited Paper
for the MAR05 Meeting of
the American Physical Society

Destabilization of light element hydrides with high hydrogen capacities: metal imides/nitrides

WEIFANG LUO

High hydrogen capacity materials are highly desirable for hydrogen storage for on-board applications. Some light elements form hydrides with high hydrogen capacities, such as LiH (12.7 wt%) and MgH₂ (7.6 wt%). These hydrides, however, are very stable, releasing hydrogen only at very high temperature, above 600°C and 350°C, respectively, with poor kinetics. These hydrogen storage features are unsatisfactory for on-board application. Chen et al [1] reported the hydrogen storage properties of lithium nitride/imide. According to their results lithium imide can absorb hydrogen at 1 bar at 285°C reversibly with hydrogen capacity of 6.5wt%. Lithium nitride, on the other hand, can absorb 5wt% more hydrogen, however, it is much more stable compared with lithium imide.



This indicates that it is an effective method to destabilize lithium hydride by converting hydride to nitrogen-containing material, such as lithium imide/nitride. Here we report a new approach to further de-stabilize lithium imide by partial substitution of lithium by magnesium in this system. This Mg-substituted material releases hydrogen of significant higher pressure at much lower temperature than those for lithium imide, with minimal capacity reduction [2]. One of the examples is the mixture of (LiNH₂-MgH₂), which can release hydrogen of approximately 30 bar at 200°C reversibly, with hydrogen capacity of 5 wt%. This material has the potential to deliver hydrogen of 3 bar at 100°C. It may be further dehydrogenated to nitride with total hydrogen capacity of approximately 9wt%. The destabilization mechanism for this system will be discussed since this may provide clue in the future searching for high capacity hydrogen storage materials [1] P. Chen, Z. Xiong, J. Luo, J. Lin, L Tan, Nature Vol. 420 (2002) 302-304. [2] W. Luo, J. Alloys and Compounds, 381 (2004) 284-287.